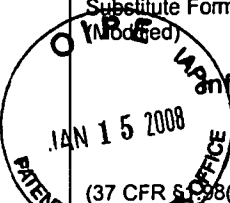


Substitute Form PTO-1449 (Modified)  Information Disclosure Statement by Applicant (Use several sheets if necessary) (37 CFR 1.128(b))	U.S. Department of Commerce Patent and Trademark Office	Attorney's Docket No. 22578-005US1	Application No. 10/535,345
	Applicant Semple, et al.		
	Filing Date February 15, 2006	Group Art Unit 1626	

## U.S. Patent Documents

Examiner Initial	Desig. ID	Document Number	Publication Date	Patentee	Class	Subclass	Filing Date If Appropriate
	AA						

## Foreign Patent Documents or Published Foreign Patent Applications

Examiner Initial	Desig. ID	Document Number	Publication Date	Country or Patent Office	Class	Subclass	Translation	
							Yes	No
	AB	WO06/069242A2	06/29/06	WIPO				
	AC	WO05044816A1	05/19/05	WIPO				
	AD	WO04103370A1	12/02/04	WIPO				
	AE	WO03078409A1	09/25/03	WIPO				
	AF	WO03062200A2	07/31/03	WIPO				
	AG	WO03022814A1	03/20/03	WIPO				
	AH	WO03002582A1	01/09/03	WIPO				
	AI	WO02094830A2	11/28/02	WIPO				
	AJ	WO0179169A2	10/25/01	WIPO				
	AK	WO0166520A1	09/13/01	WIPO				
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	AN	EP1305286B1	12/08/04	Europe				
	AO	EP05298854A2	03/09/93	Europe				

## Other Documents (include Author, Title, Date, and Place of Publication)

Examiner Initial	Desig. ID	Document
	AP	Alterman, M. et al., "Fast microwave-assisted preparation of aryl and vinyl nitriles and the corresponding tetrazoles from organo-halides", J. Org. Chem. 65:7984-89 (2000)(supporting information attached)
	AQ	Cahn, R.S. et al., "Specification of molecular chirality", Angew. Chem. Internat. Edit. 5(4):385-415 (1966)
	AR	Carballo-Jane et al., "Comparison of rat and dog models of vasodilation and lipolysis for the calculation of a therapeutic index for GPR109A agonists," <i>Journal of Pharmacological and Toxicological Methods</i> , Article in Press, doi:10.1016/j.vascn.2007.05.007 (2007).
	AS	Carballo-Jane et al., "Comparison of rat and dog models of vasodilation and lipolysis for the calculation of a therapeutic index for GPR109A agonists," <i>Journal of Pharmacological and Toxicological Methods</i> , 56(3). pp. 308-316, (2007).

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	AT	Clayton, S. et al., "A total synthesis of (±)-epibatidine", Tetrahedron Letters 34(46):7493-6 (1993)
	BA	Cohen, T. et al., "Synthetically useful $\beta$ -lithioalkoxides from reductive lithiation of epoxides by aromatic radical anions", J. Org. Chem. 55:1528-36 (1990)
	BB	Corsaro, A. et al., "Steric course of some cyclopropanation reactions of L-threo-hex-4-enopyranosides", Tetrahedron Letters 60:3787-95 (2004)
	BC	Effenberger, F. et al., "Regioselective halo- and carbodesilylation of (trimethylsilyl)-1-methylpyrazoles", J. Org. Chem. 49:4687-95 (1984)
	BD	Gharbaoui et al., "Agonist lead identification for the high affinity niacin receptor GPR109a," <i>Bioorganic &amp; Medicinal Chemistry Letters</i> , 17:4914-4919 (2007).
	BE	Hodgson, D. et al., "Intramolecular cyclopropanation of unsaturated terminal epoxides", J. Am. Chem. Soc. 126:8664-5 (2004)
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	BG	Jung et al., "Analogues of acifran: agonists of the high and low affinity niacin receptors, GPR109a and GPR109b," <i>Journal of Medicinal Chemistry</i> , 50:1445-1448 (2007).
	BH	Katritzky, A. et al., "Alpha-lithiation of N-alkyl groups in pyrazoles", Tetrahedron Letters 39:2023-9 (1983)
	BI	Latli, B. et al., "Novel and potent 6-chloro-3-pyridinyl ligands for the $\alpha$ 4 $\beta$ 2 neuronal nicotinic acetylcholine receptor", J. Med. Chem. 42:2227-34 (1999)
	BJ	Latli, B. et al., "Supporting information for "Novel and potent 6-chloro-3-pyridinyl ligands for the $\alpha$ 4 $\beta$ 2 neuronal nicotinic acetylcholine receptor", J. Med. Chem. pp. 2227(1999)
	BK	Maciejewski-Lenoir et al., "Langerhans cells release prostaglandin D <sub>2</sub> in response to nicotinic acid," <i>Journal of Investigative Dermatology</i> , 126:2637-2646 (2006).
	CA	Mahley, R. et al., "Drug therapy for hypercholesterolemia and dyslipidemia", Goodman & Gilman 36:971-1002
	CB	Mariano, P. et al., "Mechanistic aspts of gas-phase photodecarbonylation reactions of bicycle[3.1.0]hexanones", J. Org. Chem. 45:1753-62 (1980)
	CC	Miller, R.D. et al., "Deoxygenation of sulfoxides promoted by electrophilic silicon reagents: preparation of aryl-substituted sulfonium salts", J. Org. Chem. 53:5571-3 (1988)
	CD	Movassaghi, M. et al., "A direct method for the conversion of terminal epoxides into $\gamma$ -butanolides", J. Am. Chem. Soc. 124(11):2456-7 (2002)
	CE	Newman-Evans, R. et al., "The influence of intramolecular dynamics on branching ratios in thermal rearrangements", J. Org. Chem. 55:695-711 (1990)
	CF	Nishimura, J. et al., "A novel synthesis of methylcyclopropanes", Tetrahedron Letters 25:2647-59 (1969)
	CG	Olivo, H. et al., "Synthetic studies on the trans-Chlorocyclopropane diene side chain of Callipeltoside-A", Org. Lett. 2(25):4055-8 (2000)(supporting information attached)
	CH	Prelog, V. et al., "Basic principles of the CIP-system and proposals for a revision", Angew. Chem. Int. Ed. Engl. 21:567-83 (1982)
	CI	Richman et al., "Nicotinic acid receptor agonists differentially activate downstream effectors," <i>The Journal of Biological Chemistry</i> , 282:18028-18036, (2007).

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Other Documents (include Author, Title, Date, and Place of Publication)		
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	CJ	Schaus, S. et al., "Highly selective hydrolytic kinetic resolution of terminal epoxides catalyzed by chiral (salen)cobalt(III)-complexes. Practical synthesis of enantioenriched terminal epoxides and 1,2-Diols", JACS 124:1307 (2002)(supporting information attached)
	CK	Semple et al., "Recent progress in the discovery of niacin receptor agonists," <i>Current Opinion in Drug Discovery &amp; Development</i> , 10:452-459, (2007).
	CL	Semple et al., "1-Alkyl-benzotriazole-5-carboxylic acids are highly selective agonists of the human orphan G-protein-coupled receptor GPR109b," <i>Journal of Medicinal Chemistry</i> 49:1227-1230, (2006).
	CM	Semple, "Niacin receptor agonists," Presentation, American Chemical Society 233 <sup>rd</sup> National Meeting & Exposition, March 25, 2007 – March 29, 2007, Chicago, Illinois
	CN	Semple, "Discovery of selective agonists for GPR109a and GPR109b, the high and low affinity receptors for niacin," Presentation, <i>GPCRs in Medicinal Chemistry</i> , jointly organized by the Society of Chemical Industry, Royal Society of Chemistry and the Societa Chimica Italiana, September 18, 2006 – September 20, 2006, Verona, Italy
	CO	Skinner et al., "Fluorinated pyrazole acids are agonists of the high affinity niacin receptor GPR109a," Poster, 30 <sup>th</sup> National Medicinal Chemistry Symposium, June 25, 2006 – June 29, 2006, Seattle, WA
	CP	Smith, A. et al., "Total synthesis of the neotropical poison-frog alkaloid (-)-205B, Org. Lett. 7(15):3247-50 (2005)(supporting information attached)
	CQ	Taber, D. et al., "Synthesis of the eight enantiomerically pure diastereomers of the 12-F2-Isoprostanes", J. Am. Chem. Soc. 124:13121-6 (2002)(supporting information attached)
	CR	Taggart et al., "(D)-β-Hydroxybutyrate inhibits adipocyte lipolysis via the nicotinic acid receptor PUMA-G," <i>The Journal of Biological Chemistry</i> , 280:26649-26652, (2005).
	CS	Turner, S. et al., "Enantiospecific synthesis of annulated nicotine analogues from D- and L-glutamic acid Pyridotropanes", J. Org. Chem. 65:861-70 (2000)
	DA	Yagi, H. et al., "Removal of benzyl-type protecting groups from peptides by catalytic transfer hydrogenation with formic acid", J. Org. Chem. 44(19):3442-4 (1979)
	DB	Zhang, R. et al., "Cyclopropanation reactions of pyroglutamic acid-derived synthons with alkylidene transfer reagents", J. Org. Chem. 64:547-55 (1999)(supporting information attached)

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